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EXAMINER	
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	HEINCER, LIAM J ART UNIT PAPER NU

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication:

	Application No.	Applicant(s)		
	10/536,901	ONISHI, YASUHIKO		
Office Action Summary	Examiner	Art Unit		
	Liam J. Heincer	1709		
The MAILING DATE of this communication app		orrespondence address		
Period for Reply				
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tim vill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONEI	I. lely filed the mailing date of this communication. D (35 U.S.C. § 133).		
Status				
1) Responsive to communication(s) filed on 27 M	ay 2005.			
2a) This action is FINAL . 2b) ☑ This action is non-final.				
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is				
closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.				
Disposition of Claims		•		
4)⊠ Claim(s) <u>1-6</u> is/are pending in the application.				
4a) Of the above claim(s) is/are withdrawn from consideration.				
5) Claim(s) is/are allowed.				
6)⊠ Claim(s) <u>1-6</u> is/are rejected.				
7) Claim(s) is/are objected to.				
8) Claim(s) are subject to restriction and/or	relection requirement.			
Application Papers				
9)☐ The specification is objected to by the Examiner.				
10) ☐ The drawing(s) filed on is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.				
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).				
Replacement drawing sheet(s) including the correcti	, , , , ,			
11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.				
Priority under 35 U.S.C. § 119		,		
12)⊠ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a)⊠ All b)□ Some * c)□ None of:				
1. Certified copies of the priority documents have been received.				
2. Certified copies of the priority documents have been received in Application No				
3. Copies of the certified copies of the priority documents have been received in this National Stage				
application from the International Bureau (PCT Rule 17.2(a)).				
* See the attached detailed Office action for a list of the certified copies not received.				
Attachment(s)				
1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948)	4) Ll Interview Summary Paper No(s)/Mail Da			
3) Information Disclosure Statement(s) (PTO/SB/08)	5) 🔲 Notice of Informal Pa			
Paper No(s)/Mail Date <u>9/2005</u> .	6)			

DETAILED ACTION

Claims 1-4 are objected to because of the following informalities: In claims 1-4 during the description of R₇ semicolons are used to separate the members of the group rather than commas. Appropriate correction is required.

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claim 1 is rejected under 35 U.S.C. 102(b) as being anticipated by Onishi (US Pat. 4,816,540).

Considering Claim 1: Onishi teaches a cationic graft-copolymer (1:1) of a water-soluble linear backbone polymer having hydroxyl groups (2:6-10) comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)

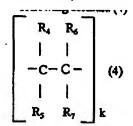
 $[C_6H_7O_2(OH)_{3-6}(OX)_a]_2H_2O(1)$

or a unit derived from a water-soluble linear polyvinyl alcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)

+CH₂ CH(OH)_{1+b} (OX)_b → (2) +CH₂ CH(OH)_{1+b} (OX)_b (OAc)_c→ (3)

Wherein X is a -- $(CH_2)_mR_1$ organic radical where R_1 is a member of the class consisting of -- NH_2 radical, -- $N(CH_3)_2$ radical, -- $N(C_2H_5)_2$ radical, -- $N^+(C_2H_5)_3$ radical, -- $N^+(CH_2)_2CH_2CH(OH)CH_3$ radical, -- $N^+(C_2H_5)_2CH_2CH(OH)CH_3$ radical, -- $N^+(C_2H_5)_2(C_2H_5)N$ ($C_2H_5)_2$ radical, -- $C_6H_4NH_2$ radical, and - $COC_6H_4NH_2$ radical, -- COR_2 radical where R_2 is -- COR_2 radical where R_3 is -- COR_2 radical where COR_3 radical value of COR_3 radical, m is a natural number of 1 to 3, a is a positive number having a value of COR_3 radical (2:13-44); and a unit derived from a polymerize-able

olefin compound of the following formula (4)



Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of

Where R_8 is a member of the class consisting of hydrogen, C_1 - C_{12} alkyl radicals, cyclohexyl radical, C_1 - C_4 hydroxyalkyl radicals, C_1 - C_8 aminoalkyl radicals, C_1 - C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1 - C_4 lower alkyl – substituted tetrahydrofuran radical, benzyl radical, the $(CH_2CH_2O)_yCH_2CH_2OH$ radical where y is a positive integer from 1 to 10,and--N(R_9)₂ where the two R_9 's which may be the same or different, are either hydrogen or a C_1 - C_4 alkyl radical;

Where R_{10} is a C_1 - C_8 alkyl radical; phenyl radical; tolyl radical; pyrrolidone radical; and

Where R₁₁ is NH₂, NHCH₃, N,N-dimethylamino radical, N,N-dimethylaminopropylamino radical, and morpholine radical (2:66-3:39). The limitation "for a non-viral gene delivery vector" is functional language that does not materially change the structure of the polymer.

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Claim 2 is rejected under 35 U.S.C. 102(b) as being anticipated by Onishi (US Pat. 4,816,540).

Considering Claim 2: Onishi teaches process for preparing a cationic graft-copolymer (1:27-31) of a water-soluble linear backbone polymer having hydroxyl groups (2:6-10) comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)

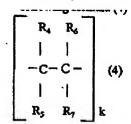
$$[C_6H_7O_2(OH)_{3-6}(OX)_a]_2H_2O(1)$$

or a unit derived from a water-soluble linear polyvinyl alcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)

$$f_{CH_2 CH(OH)_{1+}}(OX)_{b} - f_{a} (2)$$

 $f_{CH_2 CH(OH)_{1++}}(OX)_{b} (OAc)_{c} - f_{a} (3)$

Wherein X is a -- $(CH_2)_mR_1$ organic radical where R_1 is a member of the class consisting of -- NH_2 radical, -- $N(CH_3)_2$ radical, -- $N(C_2H_5)_2$ radical, -- N^+ $(C_2H_5)_3$ radical, -- N^+ $(C_2H_5)_2$ CH₂CH(OH)CH₃ radical, -- N^+ $(C_2H_5)_2$ CH₂CH(OH)CH₃ radical, -- N^+ $(C_2H_5)_2$ (C_2H_5)N $(C_2H_5)_2$ radical, -- $C_6H_4NH_2$ radical, and - $COC_6H_4NH_2$ radical, -- COR_2 radical where R_2 is -- COR_2 radical where R_3 is -- COR_2 radical where COR_3 radical, -- COR_3 radical, m is a natural number of 1 to 3, a is a positive number having a value of COR_3 radical (2:13-44); and a unit derived from a polymerize-able olefin compound of the following formula (4)



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Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of

Where R_8 is a member of the class consisting of hydrogen, C_1 - C_{12} alkyl radicals, cyclohexyl radical, C_1 - C_4 hydroxyalkyl radicals, C_1 - C_8 aminoalkyl radicals, C_1 - C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1 - C_4 lower alkyl – substituted tetrahydrofuran radical, benzyl radical, the $(CH_2CH_2O)_yCH_2CH_2OH$ radical where y is a positive integer from 1 to 10,and--N(R_9)₂ where the two R_9 's which may be the same or different, are either hydrogen or a C_1 - C_4 alkyl radical;

Where R_{10} is a C_1 - C_8 alkyl radical; phenyl radical; tolyl radical; pyrrolidone radical; and

Where R₁₁ is NH₂, NHCH₃, N,N-dimethylamino radical, N,N-dimethylaminopropylamino radical, and morpholine radical (2:66-3:39). The limitation "for a non-viral gene delivery vector" is functional language that does not materially change the structure of the polymer.

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the

invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

Claims 3 and 5 are rejected under 35 U.S.C. 103(a) as being unpatentable over Onishi (US Pat. 4,816,540) in view of Pack, Gene-Delivery Polymers.

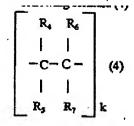
Considering Claim 3: Onishi teaches a cationic graft-copolymer (1:1) of a water-soluble linear backbone polymer having hydroxyl groups (2:6-10) comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)

$$[C_6H_7O_2(OH)_{3-4}(OX)_a]_2H_2O(1)$$

or a unit derived from a water-soluble linear polyvinyl alcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)

$$+$$
 CH₂ CH(OH)₁₋₆ (OX)₆ $+$ (2)
 $+$ CH₂ CH(OH)₁₋₆ (OX)₆ (OAc)_c $+$ (3)

Wherein X is a -- $(CH_2)_mR_1$ organic radical where R_1 is a member of the class consisting of -- NH_2 radical, -- $N(CH_3)_2$ radical, -- $N(C_2H_5)_2$ radical, -- $N^+(C_2H_5)_3$ radical, -- $N^+(CH_2)_2CH_2CH(OH)CH_3$ radical, -- $N^+(C_2H_5)_2CH_2CH(OH)CH_3$ radical, -- $N^+(C_2H_5)_2(C_2H_5)N$ $(C_2H_5)_2$ radical, -- $C_6H_4NH_2$ radical, and - $COC_6H_4NH_2$ radical, -- COR_2 radical where R_2 is -- COR_2NH_2 or -- COR_2NH_2 or -- COR_2NH_2 radical, m is a natural number of 1 to 3, a is a positive number having a value of COR_2NH_2 radical (2:13-44); and a unit derived from a polymerize-able olefin compound of the following formula (4)



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Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of

Where R_8 is a member of the class consisting of hydrogen, C_1 - C_{12} alkyl radicals, cyclohexyl radical, C_1 - C_4 hydroxyalkyl radicals, C_1 - C_8 aminoalkyl radicals, C_1 - C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1 - C_4 lower alkyl – substituted tetrahydrofuran radical, benzyl radical, the $(CH_2CH_2O)_yCH_2CH_2OH$ radical where y is a positive integer from 1 to 10,and- $N(R_9)_2$ where the two R_9 's which may be the same or different, are either hydrogen or a C_1 - C_4 alkyl radical;

Where R_{10} is a C_1 - C_8 alkyl radical; phenyl radical; tolyl radical; pyrrolidone radical; and

Where R₁₁ is NH₂, NHCH₃, N,N-dimethylamino radical, N,N-dimethylaminopropylamino radical, and morpholine radical (2:66-3:39). The limitation "for a non-viral gene delivery vector" is functional language that does not materially change the structure of the polymer.

Onishi does not teach forming a complex with DNA. However, Pack teaches forming a complex between a cationic graft polymer and DNA (Section 2.2.2). Onishi and Pack are combinable as they are concerned with the same field of endeavor, namely cationic polymers. It would have been obvious to a person having ordinary skill in the art at the time of the invention to have formed a complex between the polymer of Onishi and DNA as in Pack, and the motivation to do so would have been, as Pack suggests, to form a complex for use in gene therapy (Section 2.2).

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Considering Claim 5: Onishi does not teach a gene delivery system. However, Pack teaches forming a gene delivery system from a complex between a cationic graft polymer and DNA (Section 2.2.2). It would have been obvious to a person having ordinary skill in the art at the time of the invention to have formed the gene delivery system from the polymer of Onishi as in Pack, and the motivation to do so would have been, as Pack suggests, to treat disease with gene therapy (Section 1).

Claims 4 and 6 are rejected under 35 U.S.C. 103(a) as being unpatentable over Onishi (US Pat. 4,816,540) in view of Pack, Gene-Delivery Polymers.

Considering Claim 4: Onishi teaches a cationic graft-copolymer (1:1) of a water-soluble linear backbone polymer having hydroxyl groups (2:6-10) comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)

$$[C_6H_7O_2(OH)_{3-d}(OX)_a]_2H_2O(1)$$

or a unit derived from a water-soluble linear polyvinyl alcohol of the following formula (2) or a partial hydrolyzed alcohol of the following formula (3)

```
+CH<sub>2</sub> CH(OH)<sub>1-6</sub> (OX)<sub>6</sub> → (2)
+CH<sub>2</sub> CH(OH)<sub>1-6</sub> (OX)<sub>6</sub> (OAc)<sub>6</sub> → (3)
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olefin compound of the following formula (4)

Wherein R₄, R₅ and R₆ are each selected from the group consisting of hydrogen and CH₃ and R₇ is a member of the group consisting of

Where R₈ is a member of the class consisting of hydrogen, C₁-C₁₂ alkyl radicals, cyclohexyl radical, C₁-C₄ hydroxyalkyl radicals, C₁-C₈ aminoalkyl radicals, C₁-C₈ dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C₁-C₄ lower alkyl – substituted tetrahydrofuran radical, benzyl radical, the (CH₂CH₂O)_vCH₂CH₂OH radical where y is a positive integer from 1 to 10, and -- N(R₉)₂ where the two R₉'s which may be the same or different, are either hydrogen or a C₁-C₄ alkyl radical;

Where R₁₀ is a C₁- C₈ alkyl radical; phenyl radical; tolyl radical; pyridine radical; pyrrolidone radical; and

Where R₁₁ is NH₂, NHCH₃, N,N-dimethylamino radical, N,N-dimethylaminopropylamino radical, and morpholine radical (2:66-3:39). The limitation "for a non-viral gene delivery vector" is functional language that does not materially change the structure of the polymer.

Onishi does not teach forming a complex with RNA. However, Pack teaches forming a complex between a cationic graft polymer and RNA (Sections 2.2.2 and 2.2). Onishi and Pack are combinable as they are concerned with the same field of endeavor, namely cationic polymers. It would have been obvious to a person having ordinary skill in the art at the time of the invention to have formed a complex between the polymer of Onishi and RNA as in Pack, and the motivation to do so would have been, as Pack suggests, to form a complex for use in gene therapy (Section 2.2).

Considering Claim 6: Onishi does not teach a gene delivery system. However, Pack teaches forming a gene delivery system from a complex between a cationic graft polymer and RNA (Sections 2.2.2 and 2.2). It would have been obvious to a person having ordinary skill in the art at the time of the invention to have formed the gene delivery system from the polymer of Onishi as in Pack, and the motivation to do so would have been, as Pack suggests, to treat disease with gene therapy (Section 1).

The prior art made of record and not relied upon is considered pertinent to applicant's disclosure. See PTO form 892.

Double Patenting

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper timewise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. A nonstatutory obviousness-type double patenting rejection is appropriate where the conflicting claims are not identical, but at least one examined application claim is not patentably distinct from the reference claim(s) because the examined application claim is either anticipated by, or would have been obvious over, the reference claim(s). See, e.g., *In re Berg*, 140 F.3d 1428, 46 USPQ2d 1226 (Fed. Cir. 1998); *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422

F.2d 438, 164 USPQ 619 (CCPA 1970); and *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) or 1.321(d) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent either is shown to be commonly owned with this application, or claims an invention made as a result of activities undertaken within the scope of a joint research agreement.

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claim 1 is rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claim 1 of U.S. Patent No. 4,816,540.

Considering Claim 1: Patent '540 teaches a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)

[C₆H₇O₂(OH)₃₋₁(OX)₆]₆H₂O (1)

Wherein X is a --(CH_2)_mR₁ organic radical where R₁ is a member of the class consisting of -- NH_2 radical, -- $N(CH_3)_2$ radical, -- $N(C_2H_5)_2$ radical, -- N^+ (C_2H_5)₃ radical, --

N⁺(CH₂)₂CH₂CH(OH)CH₃ radical, -- N⁺(C₂H₅)₂CH₂CH(OH)CH₃ radical,

-- $N^{+}(C_2H_5)_2(C_2H_5)N$ ($C_2H_5)_2$ radical, -- $C_6H_4NH_2$ radical, and - $COC_6H_4NH_2$ radical, --

 COR_2 radical where R_2 is -- CH_2NH_2 or -- $C_6H_4NH_2$, -- $CH_2CH(OH)CH_2R_3$ radical where

 R_3 is --NH₂, --N(CH₃)₂, --N(C₂H₅)₂, and -N⁺(C₂H₅)₃ radical, m is a natural number of 1 to

3, a is a positive number having a value of 0<a<3, b is a positive number having a value

of 0<b<1, and Ac is acetyl radical; and a unit derived from a polymerize-able olefin

compound of the following formula (4)

Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of

Where R_8 is a member of the class consisting of hydrogen, C_1 - C_{12} alkyl radicals, cyclohexyl radical, C_1 - C_4 hydroxyalkyl radicals, C_1 - C_8 aminoalkyl radicals, C_1 - C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1 - C_4 lower alkyl – substituted tetrahydrofuran radical, benzyl radical, the $(CH_2CH_2O)_yCH_2CH_2OH$ radical where y is a positive integer from 1 to 10,and--N(R_9)₂ where the two R_9 's which may be the same or different, are either hydrogen or a C_1 - C_4 alkyl radical;

Where R_{10} is a C_1 - C_8 alkyl radical; phenyl radical; tolyl radical; pyrrolidone radical; and

Where R₁₁ is NH₂, NHCH₃, N,N-dimethylamino radical, N,N-dimethylaminopropylamino radical, and morpholine radical (Claim 1).

Claim 2 is rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claim 2 of U.S. Patent No. 4,816,540.

Considering Claim 2: Patent '540 teaches process for preparing a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)

 $[C_6H_7O_2(OH)_{3-a}(OX)_a]_xH_2O(1)$

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Wherein X is a -- $(CH_2)_mR_1$ organic radical where R_1 is a member of the class consisting of -- NH_2 radical, -- $N(CH_3)_2$ radical, -- $N(C_2H_5)_2$ radical, -- N^+ $(C_2H_5)_3$ radical, -- N^+ $(C_2H_5)_2$ CH₂CH(OH)CH₃ radical, -- N^+ $(C_2H_5)_2$ CH₂CH(OH)CH₃ radical, -- N^+ $(C_2H_5)_2$ (C_2H_5)N $(C_2H_5)_2$ radical, -- $C_6H_4NH_2$ radical, and - $COC_6H_4NH_2$ radical, -- COR_2 radical where R_2 is -- COR_2 radical where R_3 is -- COR_3 radical where COR_3 radical, -- COR_3 radical, m is a natural number of 1 to 3, a is a positive number having a value of COR_3 radical (2:13-44); and a unit derived from a polymerize-able olefin compound of the following formula (4)

Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of

Where R_8 is a member of the class consisting of hydrogen, C_1 - C_{12} alkyl radicals, cyclohexyl radical, C_1 - C_4 hydroxyalkyl radicals, C_1 - C_8 aminoalkyl radicals, C_1 - C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1 - C_4 lower alkyl – substituted tetrahydrofuran radical, benzyl radical, the $(CH_2CH_2O)_yCH_2CH_2OH$ radical where y is a positive integer from 1 to 10,and--N(R_9)₂ where the two R_9 's which may be the same or different, are either hydrogen or a C_1 - C_4 alkyl radical;

Where R₁₀ is a C₁- C₈ alkyl radical; phenyl radical; tolyl radical; pyridine radical;

pyrrolidone radical; and

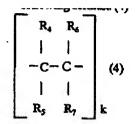
Where R₁₁ is NH₂, NHCH₃, N,N-dimethylamino radical, N,N-dimethylaminopropylamino radical, and morpholine radical (Claim 2).

Claims 3-6 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claim 1 of U.S. Patent No. 4,816,540 in view of Pack, Gene-Delivery Polymers.

<u>Considering Claims 4 and 5</u>: Patent '540 teaches a cationic graft-copolymer of a water-soluble linear backbone polymer having hydroxyl groups comprising a unit derived from a cationic water-soluble linear polysaccharide of the following formula (1)

$$[C_6H_7O_2(OH)_{3-a}(OX)_a]_1H_2O(1)$$

Wherein X is a --(CH₂)_mR₁ organic radical where R₁ is a member of the class consisting of -- NH₂ radical, -- N(CH₃)₂ radical, -- N(C₂H₅)₂ radical, -- N⁺ (C₂H₅)₃ radical, -- N⁺ (CH₂)₂CH₂CH(OH)CH₃ radical, -- N⁺ (CH₂)₂CH₂CH(OH)CH₃ radical, -- N⁺ (CH₂)₂CH₅)₂(CH₂CH₅)₂ radical, -- C₆H₄NH₂ radical, and - COC₆H₄NH₂ radical, -- COR₂ radical where R₂ is -- CH₂NH₂ or -- C₆H₄NH₂, -- CH₂CH(OH)CH₂R₃ radical where R₃ is --NH₂, --N(CH₃)₂, --N(C₂H₅)₂, and -N⁺ (C₂H₅)₃ radical, m is a natural number of 1 to 3, a is a positive number having a value of 0<a>a<3, b is a positive number having a value of 0<bar>
o



Wherein R_4 , R_5 and R_6 are each selected from the group consisting of hydrogen and CH_3 and R_7 is a member of the group consisting of

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O
||
-C-O-Ra
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Where R_8 is a member of the class consisting of hydrogen, C_1 - C_{12} alkyl radicals, cyclohexyl radical, C_1 - C_4 hydroxyalkyl radicals, C_1 - C_8 aminoalkyl radicals, C_1 - C_8 dialkylaminoalkyl radicals, glycidyl radical, tetrahydrofuran radical, C_1 - C_4 lower alkyl – substituted tetrahydrofuran radical, benzyl radical, the $(CH_2CH_2O)_yCH_2CH_2OH$ radical where y is a positive integer from 1 to 10,and- $N(R_9)_2$ where the two R_9 's which may be the same or different, are either hydrogen or a C_1 - C_4 alkyl radical;

Where R_{10} is a C_1 - C_8 alkyl radical; phenyl radical; tolyl radical; pyrrolidone radical; and

Where R₁₁ is NH₂, NHCH₃, N,N-dimethylamino radical, N,N-dimethylaminopropylamino radical, and morpholine radical (Claim 1).

Patent '540 does not teach forming a complex with DNA or RNA. However, Pack teaches forming a complex between a cationic graft polymer and DNA or RNA (Sections 2.2.2 and 2.2). Patent '540 and Pack are combinable as they are concerned with the same field of endeavor, namely cationic polymers. It would have been obvious to a person having ordinary skill in the art at the time of the invention to have formed a complex between the polymer of Patent '540 and DNA or RNA as in Pack, and the motivation to do so would have been, as Pack suggests, to form a complex for use in gene therapy (Section 2.2).

Considering Claims 5 and 6: Patent '540 does not teach a gene delivery system. However, Pack teaches forming a gene delivery system from a complex between a cationic graft polymer and DNA or RNA (Sections 2.2.2 and 2.2). It would have been obvious to a person having ordinary skill in the art at the time of the invention to have formed the gene delivery system from the polymer of Patent '540 as in Pack, and the motivation to do so would have been, as Pack suggests, to treat disease with gene therapy (Section 1).

Correspondence

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Liam J. Heincer whose telephone number is 571-270-3297. The examiner can normally be reached on Monday thru Friday 7:30 to 5:00 EST.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Mark Eashoo can be reached on 571-272-1197. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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LJH

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